

	Type	L #	Hits	Search Text	Dbs	Time Stamp	Comments	Error Definition	Error Rows
1	BRS	L1	517	(glucagon-like adj peptide) or glp-1 or glp-2	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 13:58			0
2	BRS	L2	25206	liposome	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 13:59			0
3	BRS	L3	14	1 same 2	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 14:04			0
4	BRS	L4	6	1 same conjugate	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 14:04			0

	Type	L #	Hits	Search Text	DBs	Time Stamp	Comments	Error Definition	Error Rows
1	BRS	L1	517	(glucagon-like adj peptide) or glp-1 or glp-2	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 20:14			0
2	BRS	L2	182	1 same analog	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 20:15			0
3	BRS	L3	100	1 same fragment	USPAT; US-PGPUB; EPO; DERWENT	2002/05/27 20:15			0

Glucagons and Glucagon-Like Peptides

-15 °C	Glucagon (1-29) (human, bovine, porcine)	H-6790.0500	0.5 mg	180.-
	H-His-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Lys-Tyr-Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-OH $C_{151}H_{222}N_{40}O_{45}S$ M _r :3482.80 [16941-32-5]	H-6790.1000	1 mg	320.-
-15 °C	(Des-His¹ Glu⁹)-Glucagon (1-29) amide (human, bovine, porcine)	H-2754.0500	0.5 mg	135.-
	H-His-Gln-Gly-Thr-Phe-Thr-Ser-Glu-Tyr-Ser-Lys-Tyr-Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-NH ₂ $C_{148}H_{211}N_{40}O_{45}S$ M _r :3358.70 [110084-95-2]	H-2754.1000	1 mg	200.-
Glucagon antagonist: Lit. C.G.Unson et al., Peptides 10 , 1171 (1989)				
-15 °C	Glucagon (1-37) (porcine)	H-6880.0500	0.5 mg	430.-
	H-His-Ser-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Lys-Tyr-Leu-Asp-Ser-Arg-Arg-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-Lys-Arg-Asn-Lys-Asn-Ile-Ala-OH (Oxytomodulin (porcine)) Lit. D.Boraille et al., Peptides 2 (suppl. 2), 41 (1981) / D.Boraille et al., Ann. N.Y. Acad. Sci. 527 , 168 (1988)	H-6880.1000	1 mg	785.-
-15 °C	Glucagon (19-29) (human, bovine, porcine)	H-2758.0001	1 mg	35.-
	H-Ala-Gln-Asp-Phe-Val-Gln-Trp-Leu-Met-Asn-Thr-OH $C_{61}H_{99}N_{13}O_{18}S$ M _r :1352.53 [64790-15-4]	H-2758.0005	5 mg	145.-
This glucagon fragment inhibited both the Ca ²⁺ activated and Mg ²⁺ dependent ATPase activity and Ca ²⁺ transport in liver plasma membranes with an efficiency 1000-fold higher than that of glucagon. It is likely to be the active peptide involved in the inhibition of the liver Ca ²⁺ pump. Lit. A.Mollat et al., Nature 325 , 620 (1987)				
-15 °C	Glucagon-Like Peptide 1 amide (human)	H-6025.0500	0.5 mg	215.-
	H-His-Asp-Glu-Phe-Glu-Arg-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Tyr-Tyr-Leu-Glu-Gln-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-NH ₂ (GLP-1 amide (human); Preproglucagon (72-107) amide (human)) $C_{142}H_{217}N_{31}O_{37}S$ M _r :4111.50 [99658-04-5]	H-6025.1000	1 mg	395.-
Lit. D.J.Drucker et al., Proc. Natl. Acad. Sci. USA 84 , 3434 (1987) / J.J.Holst et al., FEBS Lett. 211 , 169 (1987)				

-15 °C

Glucagon-Like Peptide 1 (7-36) amide (human)H-6795.0500 0.5 mg 180.-
H-6795.1000 1 mg 320.-H-His-Ala-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-NH₂
(GLP-1 (7-36) amide (human); Preproglucagon (78-107) amide (human))
 $C_{142}H_{217}N_{31}O_{37}S$ M_r:3297.68 [107444-51-9]

This GLP-1 fragment is secreted from the lower small intestine and shows a strong insulinotropic effect. It is presently considered as the most important incretin hormone. Its action is mediated by receptors expressed by the endocrine pancreatic B-cells.

Lit. B.Kaymann et al., Lancet **1987 II**, 1300 / G.I.Bell et al., Nature **304**, 368 (1983) / C.Ostkov and J.H.Nielsen, FEBS Lett. **229**, 175 (1988) / J.P.Raufman et al., J. Biol. Chem. **267**, 21432 (1992) / P.A.Martin and A.Faulker, Comp. Biochem. Biophys. **105A**, 705 (1993) / H.C.Fehmman et al., Peptides **15**, 453 (1994) / M.A.Nauck et al., Exp. Clin. Endocrinol. Diabetes **105**, 187 (1997)

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new (Ser⁷)-Glucagon-Like Peptide 1 (7-36) amide (human)H-4592.0500 0.5 mg 180.-
H-4592.1000 1 mg 320.-H-His-Ser-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Lys-Glu-Phe-Ile-Ala-Trp-Leu-Val-Lys-Gly-Arg-NH₂
(Ser⁷)-GLP-1 (7-36) amide (human); (Ser⁷)-Prepro-glucagon (78-107) amide (human))The replacement of alanine by serine significantly improved the plasma stability of GLP-1 (7-36) amide against DPP IV without impairing its insulinotropic activity. This may indicate that this modification could improve the potential of GLP-1 in the treatment of type-II diabetes.
Lit. U.Ritzel et al., J. Endocrinol. **159**, 93 (1998)**new Glucagon-Like Peptide 2 (human)**H-4766.0001 1 mg 255.-
H-4766.0500 0.5 mg 145.-H-His-Ala-Asp-Gly-Ser-Phe-Ser-Asp-Glu-Met-Asn-Thr-Ile-Leu-Asp-Asn-Leu-Ala-Ala-Lys-Phe-Ile-Asn-Trp-Leu-Gln-Thr-Lys-Ile-Thr-Asp-Arg-OH
(GLP-2 (human); Preproglucagon (126-159) (human))
 $C_{171}H_{246}N_{40}O_{45}S$ M_r:3922.35Like GLP-1, GLP-2 is secreted from enteroendocrine cells in a nutrient-dependent manner in both rodents and humans. Currently GLP-2 is used as a potential therapeutic agent for human subjects with a broad variety of intestinal diseases characterized by intestinal damage and insufficiency.
Lit. D.J.Drucker, Trends Endocrinol. Metab. **10**, 153 (1999) / D.J.Drucker et al., J. Parenter. Nutr. **23**, 598 (1999)**-15 °C D-Gluconyl-Val-Leu-Gly-Lys-NHEt**H-8530.0001 1 mg 30.-
H-8530.0005 5 mg 115.-
H-8530.0025 25 mg 455.- $C_{27}H_{51}N_5O_{16}$ M_r:620.74 [121459-49-2]
Inhibitor of the Plasmodium falciparum proteinase (K_i = 480 μM) and of the erythrocyte invasion by Plasmodium falciparum merozoites (IC₅₀ = 900 μM).
Lit. R.Mayer et al., J. Med. Chem. **34**, 3029 (1991)**Glucose-Dependent Insulinotropic Polypeptide (human)**H-5645.0500 0.5 mg 215.-
H-5645.1000 1 mg 395.-

(See Gastric Inhibitory Polypeptides and Fragments Page 415)

(H-Glu(Cys-βNA)-OH)₂ (Disulfide bond)K-1650.0050 50 mg 70.-
K-1650.0250 250 mg 285.- $C_{68}H_{106}N_8O_{25}S_2$ M_r:748.88